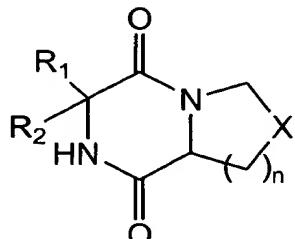


12. (Amended twice) A method of providing neuroprotection to a subject comprising administering to a subject in need of such treatment an effective amount of a compound having the formula:

1 P E



or a pharmaceutically acceptable salt or hydrate thereof, wherein:

n is an integer from 0 to 3;

X is selected from the group consisting of -S-, -O-, -NR- and -CH2-;

R<sub>1</sub> and R<sub>2</sub> are each independently selected from the group consisting of -H, -OR, -SR, -NRR, -NO<sub>2</sub>, -CN, -C(O)OR, -C(O)NRR, -C(NR)NRR, trihalomethyl, halogen, (C<sub>1</sub>-C<sub>6</sub>) alkyl, substituted (C<sub>1</sub>-C<sub>6</sub>) alkyl, (C<sub>2</sub>-C<sub>6</sub>) alkenyl, substituted (C<sub>2</sub>-C<sub>6</sub>) (C<sub>2</sub>-C<sub>6</sub>) alkenyl, (C<sub>2</sub>-C<sub>6</sub>) alkynyl, substituted (C<sub>2</sub>-C<sub>6</sub>) alkynyl, (C<sub>5</sub>-C<sub>20</sub>) aryl, substituted (C<sub>5</sub>-C<sub>20</sub>) aryl, 5-20 membered heteroaryl, substituted 5-20 membered heteroaryl, (C<sub>6</sub>-C<sub>26</sub>) alkaryl, substituted (C<sub>6</sub>-C<sub>26</sub>) alkaryl, 6-26 membered alk-heteroaryl and substituted 6-26 membered alk-heteroaryl,

or R<sub>1</sub> and R<sub>2</sub> taken together are -CH<sub>2</sub>-(CH<sub>2</sub>)<sub>m</sub>-CH<sub>2</sub>-, where m is an integer from 0 to 6;

each alkyl, alkenyl, alkynyl, aryl, alkaryl, heteroaryl or alk-heteroaryl substituent is independently selected from the group consisting of -OR, -SR, -NRR, -CN, -NO<sub>2</sub>, -C(O)OR, -C(O)NRR, -C(S)NRR, -C(NR)NRR, halogen and trihalomethyl; and

each R is independently selected from the group consisting of -H, (C<sub>1</sub>-C<sub>6</sub>) alkyl, (C<sub>2</sub>-C<sub>6</sub>) alkenyl, (C<sub>2</sub>-C<sub>6</sub>) alkynyl, (C<sub>5</sub>-C<sub>20</sub>) aryl, 5-20 membered heteroaryl, (C<sub>6</sub>-C<sub>26</sub>) alkaryl and 6-26 membered alk-heteroaryl.